

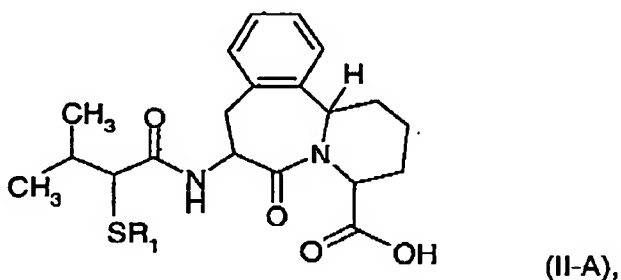
**AMENDMENTS TO THE CLAIMS**

This listing of the claims will replace all prior versions including the claims in the application.

Listing of the claims:

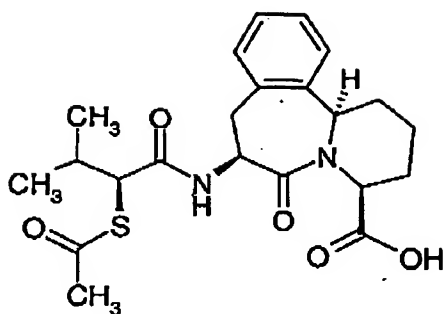
1. (Cancelled)
2. (Currently amended) The method according to claim 311 wherein the disease is selected from the group consisting of ~~non-diabetic nephropathy~~, diabetic nephropathy, insulin resistance, diabetic neuropathy, diabetic retinopathy, myocardial infarction, cataracts and diabetic cardiomyopathy.
3. (Cancelled)
4. (Original) The method according to claim 2 wherein the disease is diabetic nephropathy.
5. (Original) The method according to claim 2 wherein the disease is insulin resistance.
6. (Original) The method according to claim 2 wherein the disease is diabetic neuropathy.
7. (Original) The method according to claim 2 wherein the disease is diabetic retinopathy.
8. (Original) The method according to claim 2 wherein the disease is myocardial infarction.
9. (Original) The method according to claim 2 wherein the disease is cataracts.

10. (Original) The method according to claim 2 wherein the disease is diabetic cardiomyopathy.
11. – 13. (Cancelled)
14. (Currently Amended) The method according to claim 31~~43~~, wherein R<sub>1</sub> is acetyl.
15. (Currently Amended) The method according to claim 31~~43~~, wherein R<sub>1</sub> is hydrogen.
16. (Currently Amended) The method according to claim 31~~43~~, wherein B<sub>1</sub> and B<sub>2</sub> are hydrogen.
17. (Currently Amended) The method according to claim 31~~43~~, wherein X is – CH<sub>2</sub>.
18. (Currently Amended) The method according to claim 31~~4~~, wherein the compound is the compound of formula (II-A)



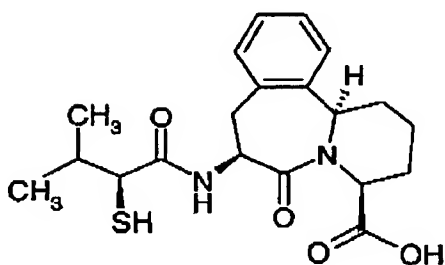
wherein R<sub>1</sub> is acetyl or hydrogen.

19. (Original) The method according to claim 18, wherein the compound has the formula (II-B)



(II-B).

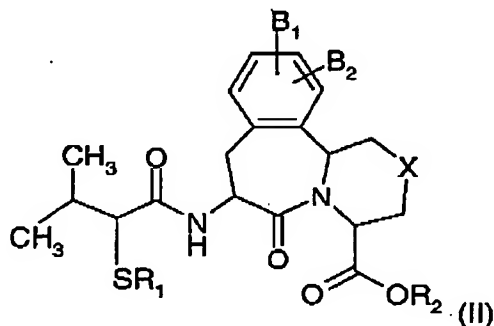
20. (Original) The method according to claim 18, wherein the compound has the formula (II-C)



(II-C).

21.-30. (Cancelled)

31. (New) A method of inhibiting both angiotensin converting enzyme and neutral endopeptidase for treatment of a disease amenable to treatment with a compound that inhibits both angiotensin converting enzyme and neutral endopeptidase which comprises administering to a patient in need of said treatment a therapeutically effective amount of a compound of formula (II)

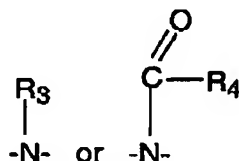


wherein

$R_1$  is hydrogen or acetyl;

$R_2$  is hydrogen,  $-\text{CH}_2\text{O}-\text{C}(\text{O})\text{C}(\text{CH}_3)_3$ ,  $\text{C}_1\text{-C}_4\text{-alkyl}$ , aryl,  $-(\text{C}_1\text{-C}_4\text{-alkyl})\text{-aryl}$ , or diphenylmethyl;

X is  $-(\text{CH}_2)_n$  wherein n is an integer 0 or 1,  $-\text{S}-$ ,  $-\text{O}-$ ,



wherein  $R_3$  is hydrogen,  $\text{C}_1\text{-C}_4\text{-alkyl}$ , aryl, or  $-(\text{C}_1\text{-C}_4\text{-alkyl})\text{-aryl}$ ; and  $R_4$  is  $\text{CF}_3$ ,  $\text{C}_1\text{-C}_{10}\text{-alkyl}$ , aryl, or  $-(\text{C}_1\text{-C}_4\text{-alkyl})\text{-aryl}$ ;

$B_1$  and  $B_2$  are each independently hydrogen, hydroxy, or  $-\text{OR}_5$ , wherein  $R_5$  is  $\text{C}_1\text{-C}_4\text{-alkyl}$ , aryl, or  $-(\text{C}_1\text{-C}_4\text{-alkyl})\text{-aryl}$  or, where  $B_1$  and  $B_2$  are attached to adjacent carbon atoms,  $B_1$  and  $B_2$  can be taken together with said adjacent carbon atoms to form a benzene ring or methylenedioxy,

or a pharmaceutically acceptable salt or stereoisomer thereof.